

What is claimed:

1. A co-formulation or kit comprising:
 - (a) a pharmaceutically effective dosage of one or more a glucose-level-controlling bioactive agents selected from an α -glucosidase inhibitor, sulfonylurea, meglitinide, thiazolidinediones, biguanide, insulin, dual PPAR α/γ agonist, PPAR γ agonist or insulin secretagogue; and
 - (b) a pharmaceutically effective dosage of (i) one or more of an antihypertensive bioactive agent selected from an ACE inhibitor, calcium channel blocker, beta blocker, angiotension II receptor antagonist or diuretic, or (ii) one or more of an anti-dyslipidemia bioactive agent selected from a HMG-CoA reductase inhibitor, bile acid sequestrant, fibric acid derivative, sterol, cholesterol absorption inhibitor, MTP inhibitor or nicotinic acid derivative;

wherein:

in the case of (i) a combination of a first bioactive agent of group (a) that is metformin with a second bioactive agent of group (b), or (ii) a combination of a first bioactive agent of group (a) that is a thiazolidinedione or dual PPAR α/γ agonist with an angiotension II receptor antagonist, one or more of the following applies:

- (I) one of the first bioactive agent or the second bioactive agent is formulated for sustained release, and the other is formulated for immediate release, each formulated for once-a-day dosing; or
- (II) the co-formulation or kit comprises (A) a biguanide and a thiazolidinedione and (B) one or more group (b) bioactive agents.

2. The kit of claim 1.
3. The co-formulation of claim 1, wherein (I) applies.
4. The co-formulation of claim 3, wherein the first bioactive agent is of group (a), and the second bioactive agent is of group (b).

5. The co-formulation of claim 4, comprising a biguanide formulated for sustained release.
6. The co-formulation of claim 5, wherein the biguanide is metformin.
7. The co-formulation of claim 5, comprising a statin.
8. The co-formulation of claim 5, comprising a thiazolidinedione.
9. The co-formulation of claim 8, comprising a statin.
10. The co-formulation of claim 8, comprising an ACE inhibitor or an angiotensin II receptor antagonist.
11. The co-formulation of claim 10, comprising a statin.
12. The co-formulation of claim 8, comprising a calcium channel blocker.
13. The co-formulation of claim 12, comprising a statin.
14. The co-formulation of claim 1, comprising a capsule wherein one or more group (a) bioactive agents are formulated in sustained release beads comprised within the capsule; and one or more group (b) bioactive agents in a more immediate release form are comprised within the capsule.
15. The co-formulation of claim 14, comprising a biguanide formulated in the sustained release beads.
16. The co-formulation of claim 15, wherein the biguanide is metformin.
17. The co-formulation of claim 15, comprising a statin.

18. The co-formulation of claim 15, comprising a thiazolidinedione formulated for sustained release.
19. The co-formulation of claim 18, comprising a statin.
20. The co-formulation of claim 18, comprising an ACE inhibitor or an angiotensin II receptor antagonist.
21. The co-formulation of claim 20, comprising a statin.
22. The co-formulation of claim 18, comprising a calcium channel blocker.
23. The co-formulation of claim 22, comprising a statin.
24. The co-formulation of claim 14, wherein the immediate release form of group (b) bioactive agent(s) is comprised of a coating on the beads.
25. The co-formulation of claim 1, comprising a compression formulation wherein
one or more group (a) bioactive agents are formulated in sustained release form
comprised within a portion of the compression formulation; and
one or more group (b) bioactive agents in a more immediate release form are comprised
within another portion of the compression formulation.
26. The co-formulation of claim 25, comprising a biguanide formulated for sustained release.
27. The co-formulation of claim 26, wherein the biguanide is metformin.
28. The co-formulation of claim 26, comprising a statin.

29. The co-formulation of claim 26, comprising a thiazolidinedione formulated for sustained release.
30. The co-formulation of claim 29, comprising a statin.
31. The co-formulation of claim 29, comprising an ACE inhibitor or an angiotensin II receptor antagonist.
32. The co-formulation of claim 31, comprising a statin.
33. The co-formulation of claim 29, comprising a calcium channel blocker.
34. The co-formulation of claim 33, comprising a statin.
35. The co-formulation of claim 1, comprising a suspension formulation wherein
one or more group (a) bioactive agents are formulated in sustained release form
comprised within particles that are suspended or adapted to be suspended in a liquid;
and
one or more group (b) bioactive agents are dissolved in the liquid.
36. The co-formulation of claim 35, comprising a biguanide formulated for sustained release.
37. The co-formulation of claim 36, wherein the biguanide is metformin.
38. The co-formulation of claim 36, comprising a statin.
39. The co-formulation of claim 36, comprising a thiazolidinedione formulated for sustained release.
40. The co-formulation of claim 39, comprising a statin.

41. The co-formulation of claim 39, comprising an ACE inhibitor or an angiotensin II receptor antagonist.
42. The co-formulation of claim 41, comprising a statin.
43. The co-formulation of claim 39, comprising a calcium channel blocker.
44. The co-formulation of claim 43, comprising a statin.
45. The co-formulation of claim 1, wherein one or more of the group (a) bioactive agents is a sulfonylurea, meglitinide, thiazolidinedione, biguanide or PPAR γ agonist.
46. The co-formulation of claim 45, wherein one or more of the group (a) bioactive agents is Glimepiride, Glipizide, Repaglinide, Pioglitazone, Rosiglitazone, Troglitazone or Metformin.
47. The co-formulation of claim 1, wherein one or more of the group (b) bioactive agents is a HMG-CoA reductase inhibitor, fibric acid derivative or MTP inhibitor
48. The co-formulation of claim 47, wherein one or more of the group (b) bioactive agents is Atorvastatin, Cerivastatin, Fluvastatin, Lovastatin, Pravastatin, Rosuvastatin, Simvastatin, Clofibrate, Fenofibrate, Febfirbozil, Ciprofibrate or Bezafibrate.
49. The co-formulation of claim 1, wherein one or more of the group (b) bioactive agents is a ACE inhibitor that is Captopril, Enalapril, Lisinopril or Ramipril.
50. The co-formulation of claim 1, wherein one or more of the group (b) bioactive agents is a calcium channel blocker that is Amlodipine, Felodipine, Nifedipine or Verapamil.

51. The co-formulation of claim 1, wherein one or more of the group (b) bioactive agents is an angiotension II receptor antagonist that is Irbesartan, Losartan or Valsartan.
52. A method of treating diabetes comprising administering a co-formulation of claim 1.
53. A method for delivering in the co-formulation a glucose-level-controlling bioactive agent and a second bioactive agent for treating a co-morbidity of diabetes, the glucose-level-controlling bioactive agent having a first dosing regimen and the second bioactive agent having a second, distinct dosing regimen, wherein the co-formulation provides a pharmacokinetic profile of the glucose-level-controlling bioactive agent that mimics the first dosing regimen and a pharmacokinetic profile of the second bioactive agent that mimics the second dosing regimen.